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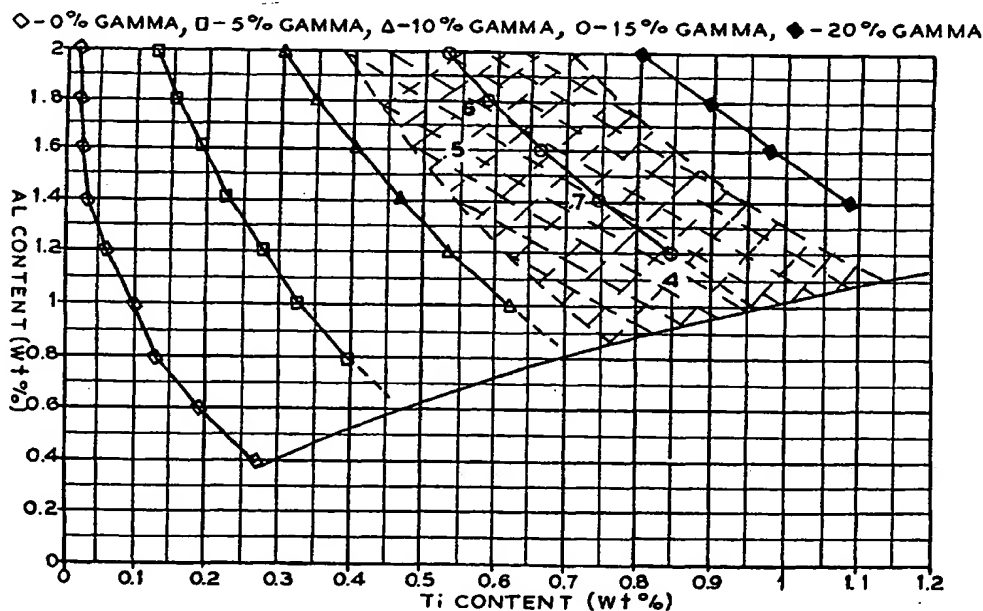
(57) Abstract

An ultra supercritical boiler tubing alloy characterized by a microstructure stabilized and strengthened for 375 bar/700 °C steam, in weight percent, service and alloyed to resist coal ash/flue gas corrosion for 200,000 hours consisting of 10 to 24 cobalt, 22.6 to 30 chromium, 2.4 to 6 molybdenum, 0 to 9 iron, 0.2 to 3.2 aluminum, 0.2 to 2.8 titanium, 0.1 to 2.5 niobium, 0 to 2 manganese, 0 to 1 silicon, 0.01 to 0.3 zirconium, 0.001 to 0.01 boron, 0.005 to 0.3 carbon, 0 to 4 tungsten, 0 to 1 tantalum and balance nickel and incidental impurities.

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ADVANCED ULTRA-SUPERCRITICAL BOILER TUBING ALLOY

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FIELD OF THE INVENTION

This invention relates to high strength alloys that possess corrosion resistance at high temperatures. In particular, it's related to alloys having sufficient strength, corrosion resistance,

15   thermal stability and fabricability for service as ultra supercritical boiler tubing.

BACKGROUND OF THE INVENTION

To meet the ever-increasing demand for electric power at lower consumer cost, utility

20   plant designers are planning new facilities that will operate at steam pressures and temperatures well beyond those currently employed. To meet their target efficiencies of greater than 60% versus today's design efficiencies of around 47%, steam parameters must be raised to 375 bar/700°C from today's typical 290 bar/580°C conditions. Unfortunately, an economical superheater tubing alloy capable of meeting the advanced steam parameters does not currently

25   exist. Examination of currently available boiler tube alloys reveals the difficulty in achieving the required combination of strength, corrosion resistance, thermal stability and manufacturability.

To meet the 38 MPa (developed by the 375 bar steam pressure) hoop stress requirement of the steam tubing, the 100,000 hour stress rupture life must exceed 100 MPa at 750°C (midradius tube wall temperature needed to maintain a 700°C steam temperature at the inner wall

5   surface). Because coal ash corrosion is capable of degrading most superheater tubing alloys, utility plant designers have placed stringent corrosion allowances on candidate alloys. The total metal loss due to inner wall steam erosion and outer wall coal ash/flue gas corrosion must not exceed 2 mm in 200,000 hours (target life for the superheater section of the steam boiler operating between 700°C and 800°C). In addition, the tube size, primarily for purposes of

10   economy, must not exceed 50 mm outer diameter (O.D.) and 8 mm wall thickness and optimally be less than 40 mm O.D. with a maximum wall thickness of 6 mm. Further, the alloy must be fabricable in high yield using conventional tube-making practices and equipment. This places a maximum constraint on work-hardening rate and yield strength of the candidate alloy range, which runs counter to the need for superior strength and stress rupture life at service

15   temperatures.

To achieve the boiler tube strength requirement, ferritic and austenitic steels must be excluded and even nickel-base solid solution alloys lack adequate strength. One is required to consider gamma prime containing alloys even though virtually all gamma prime containing alloys

20   lack adequate chromium to ensure satisfactory coal ash/flue gas corrosion resistance and few of these high strength alloys possess adequate formability to be made into tubing. Adding chromium can degrade the strengthening mechanism and if added in excess, depending on composition, can result in embrittling sigma, mu or alpha-chromium precipitation. To ensure acceptable resistance on the steam side to "downtime" stress corrosion cracking (SCC) and

25   pitting beneath the scale during operation, a minimum nickel and, advantageously a minimum molybdenum content must be present in the alloy. Since 700°C to 800°C is a very active range for carbide formation and embrittling phase precipitation, alloying content in the nickel plus cobalt matrix must be precisely limited.

30   Field fabrication places further restraint on composition. Tubing must be amenable to bending and welding during superheater boiler construction, necessitating that the alloy be supplied in the annealed condition (lowest possible practical yield strength). Whereas the strength requirement during operation demands the highest possible strength at operating temperature, requiring that the alloy be given an aging heat treatment to achieve peak strength

35   values. Dealing with these seemingly incongruous constraints and ultimately resolving this material challenge with a cost effective alloy is the object of this invention.

SUMMARY OF THE INVENTION

An ultra supercritical boiler tubing alloy characterized by a microstructure stabilized and strengthened for 375 bar/700°C steam, service and alloyed to resist coal ash/flue gas corrosion for 200,000 hours consisting of, in weight percent, 10 to 24 cobalt, 22.6 to 30
5   chromium, 2.4 to 6 molybdenum, 0 to 9 iron, 0.2 to 3.2 aluminum, 0.2 to 2.8 titanium, 0.1 to 2.5 niobium, 0 to 2 manganese, 0 to 1 silicon, 0.01 to 0.3 zirconium, 0.001 to 0.01 boron, 0.005 to 0.3 carbon, 0 to 4 tungsten, 0 to 1 tantalum and balance nickel and incidental impurities.

DESCRIPTION OF THE DRAWING

10   Figure 1 plots time to failure (logarithmic) versus temperature at an isostress of 100 MPa.

Figure 2 depicts volume gamma prime from 0 to 20 volume percent for nickel-20 cobalt-23 chromium-3 molybdenum as a function of the aluminum, titanium and niobium contents at a
15   titanium to niobium ratio of 1.5 to 0.7.

Figure 3 depicts the compositional phase boundaries at 750°C for the formation of sigma phase for the aluminum range 0.6 to 1.2 weight percent in a nickel-20 cobalt-23 chromium-6 molybdenum matrix.
20

DESCRIPTION OF PREFERRED EMBODIMENT

This Ni-Cr-Co alloy has sufficient strength, corrosion resistance, thermal stability and fabricability to serve several demanding high temperature applications. Its high chromium in combination with a relatively small amount of molybdenum and gamma prime strengtheners
25   increase corrosion resistance and strength of the alloy without compromising the thermal stability and fabricability of the alloy.

Functions of several critical elements determine the alloys' properties or characteristics. For example, the combined elements aluminum, niobium and titanium define the minimum
30   strength and limit the fabricability of the alloy. In addition, the total proportion of chromium, aluminum, titanium and niobium, as given in Equation (1), defines the minimum compositional limits for acceptable resistance to coal ash/flue gas corrosion. Carbide and embrittling phase formation chiefly limit the maximum content of the weight percent of chromium, molybdenum, titanium, niobium and carbon. See Equation (2). A minimum of 45 weight percent nickel and
35   2.4 weight percent molybdenum provide resistance to steam side stress corrosion cracking (SCC) and pitting attack – this specification refers to all elements in weight percent, unless expressly stated otherwise. A narrow range of weight percent aluminum, titanium and niobium facilitate

field fabricability and aging heat treatments such that the alloy will age to peak strength in the first few hours of operation, yet possess adequate field fabricability. The minimum alloying content for acceptable stress rupture strength is given by Equation (3) and the maximum alloying content for acceptable field fabricability by Equation (4).

5

The maximum metal loss by corrosion of 2 mm in 200,000 hours at 700°C to 800°C is achieved when the compositional limit is:

$$\%Cr + 0.6 \times \%Ti + 0.5 \times \%Al + 0.3 \times \%Nb \geq 24\%, \quad \text{preferably} \geq 25\% \qquad (1)$$

10

   Note: Equation 1 requires a minimum weight percent chromium equal to at least
           22.6% for adequate corrosion resistance.

While the maximum compositional limit below which embrittling phase precipitation

15   tends to be absent is:

$$\%(Cr + 0.8\% \times Mo) + 0.6 \times \%Ti + 0.5 \times \%Al + 0.3 \times \%Nb \leq 37.5\%, \text{preferably} \leq 35.0\% \quad (2)$$

   Note: Equation 2 defines a maximum weight percent chromium of less than or
20   equal to 30, preferably 28.5 to limit detrimental embrittling phases such as
           sigma. To minimize carbide precipitation, the maximum weight percent carbon
           must be less than or equal to 0.30, and preferably less than 0.15.

The formula for minimum aged stress rupture strength of 100,000 hours at 750°C and

25   100 MPa is:

$$\%Al + 0.56 \times \%Ti + 0.29x \, \%Nb \geq 1.7\%, \quad \text{preferably} \geq 2.0\%. \qquad (3)$$

While the maximum compositional limit to ensure fabricability is:

30

$$\%Al + 0.56 \times \%Ti + 0.29 \times \%Nb \leq 3.4\%, \quad \text{preferably} \leq 3.0\%. \qquad (4)$$

A minimum of at least 10 weight percent cobalt strengthens the matrix and increases stress rupture strength. Further increases in cobalt facilitate achieving the requisite strength.

35   Cobalt levels above 24 weight percent however, tend to diminish the thermal stability of the alloy. The alloy accepts iron as an impurity. Generally, decreasing iron content, improves

corrosion resistance properties. But for commercial recycling concerns, allowing small amounts of iron into the alloy facilitates lowering the cost of this alloy.

The alloy tolerates up to 2 weight percent manganese without an excess loss in corrosion
5   properties. Although not critical for the alloy, this element contributes to fabricability and strength by tying up sulfur. In addition, this alloy system accepts silicon as an impurity. For commercial cost considerations however, the alloy may contain up to one weight percent silicon.

As little as 0.01 weight percent zirconium improves the oxidation resistance of the alloy.
10   Zirconium in excess of 0.3 weight percent however decreases the fabricability of the alloy. Unlike zirconium, 0.001 to 0.01 weight percent boron enhances the hot workability of the alloy. Carbon in amounts of 0.005 to 0.3 weight percent provides further strength to the matrix.

Tantalum and tungsten represent important impurities that detract from the overall
15   properties of the alloy. To maintain corrosion resistance and thermal stability, total molybdenum plus tantalum plus tungsten should remain below 6 weight percent. For optimum results, this total remains below 3.5 weight percent.

Equations 1 to 4, in conjunction with Table 1 below, define "about" the ranges for
20   achieving the best combination of properties.

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TABLE 1

ELEMENT	BROAD RANGE (wt%)		INTERMEDIATE RANGE (wt%)		NARROW RANGE (wt%)		NOMINAL RANGE (wt%)	
Ni	Bal.[1]		Bal.[1]		Bal.[1]		Bal.[1]	
Co	10	- 24	12	- 23	15	- 22	18	- 21
Cr	22.6	- 30	22.8	- 29	23	- 28	23.5	- 25
Mo	2.4	- 6	2.4	- 5	2.5	- 3.5	2.8	- 3.2
Fe	0	- 9	0.1	- 4	0.3	- 2	0.5	- 1
Al	0.2	- 3.2	0.3	- 2.5	0.5	- 2	0.8	- 1.8
Ti	0.2	- 2.8	0.3	- 2.7	0.4	- 2.6	0.5	- 2.5
Nb	0.1	- 2.5	0.2	- 2.2	0.8	- 2.1	1.2	- 2
Mn	0	- 2	0.1	- 1	0.1	- 0.5	0.2	- 0.4
Si	0	- 1	0.1	- 0.8	0.2	- 0.7	0.3	- 0.6
Zr	0.01	- 0.3	0.01	- 0.2	0.03	- 0.15	0.05	- 0.1
B	0.001	- 0.01	0.002	- 0.009	0.003	- 0.008	0.004	- 0.007
C	0.005	- 0.3	0.01	- 0.2	0.02	- 0.15	0.03	- 0.12
W	0	- 4	0	- 2	0	- 1	0	- 0.8
Ta	0	- 1	0	- 0.8	0	- 0.7	0	- 0.5

(1) Plus incidental impurities

Table 2A provides specific examples of alloys within the scope of the invention.

5

TABLE 2A

ALLOY	C	Si	Mn	Fe	Ni	Cr	Ti	Al	Co	Mo	Nb
1	0.07	0.33	0.26	0.62	45.15	23.04	2.03	0.59	19.79	5.89	2.05
2	0.08	0.34	0.35	0.61	44.77	22.93	2.16	0.80	19.34	5.88	2.06
3	0.04	0.31	0.31	0.65	49.28	22.93	1.82	0.87	19.89	3.09	0.70
4	0.03	0.30	0.31	0.68	48.77	22.95	0.85	1.10	19.86	3.09	1.94
5	0.03	0.33	0.31	0.63	48.07	24.07	0.53	1.59	19.89	3.10	1.32
6	0.03	0.34	0.31	0.61	47.72	24.01	0.57	1.79	19.92	3.08	1.42
7	0.03	0.33	0.31	0.61	47.76	24.01	0.71	1.41	19.91	3.08	1.71

10   All of the compositions contained in Table 2A nominally contained 0.005 weight percent

boron, 0.06 to 0.15 weight percent zirconium, less than 0.05 weight percent tantalum and less

than 0.2 weight percent tungsten.

        Table 2B provides specific examples of alloys outside of the scope of this invention.

15

TABLE 2B

ALLOY	C	Si	Mn	Fe	Ni	Cr	Ti	Al	Co	Mo	Nb	OTHER
A	0.06	0.50	0.30	0.70	74.24	20.00	2.30	1.20	–	–	0.70	–
B	0.13	1.00	1.00	1.50	54.37	20.00	2.50	1.50	18.00	–	–	–
C	0.12	1.00	1.00	1.00	81.10	15.00	1.20	4.70	–	5.00	–	
D	0.07	0.50	0.06	–	46.23	28.50	2.35	1.35	20.3	–	0.64	–
E	0.08	0.50	1.00	7.00	72.32	15.00	2.50	0.70	–	–	0.90	–
F	0.03	0.35	0.35	18.20	52.67	19.00	0.90	0.50	–	3.00	5.00	–
G	0.03	0.50	0.50	9.50	59.47	30.00	–	–	–	–	–	–
H	0.08	0.50	0.50	1.50	52.92	22.00	0.30	1.20	12.50	9.00	–	–
I	0.05	–	–	0.50	51.10	48.00	0.35	–	–	–	–	–
J	0.55	0.40	1.00	31.75	20.00	22.00	–	0.20	18.00	3.00	–	W 2.50 Ta 0.60
K	0.05	0.56	0.22	8.76	34.82	29.99	0.25	3.06	19.97	0.29	1.89	–
L	0.03	0.31	0.42	1.93	48.99	22.77	1.18	0.62	19.40	3.02	1.19	–
M	0.09	0.83	0.21	1.55	47.41	27.99	1.53	0.21	19.97	0.12	0.01	–

SCREENING FOR COAL ASH/FLUE GAS CORROSION RESISTANCE

It is imperative that a candidate superheater boiler tubing alloy exhibit superior coal ash/flue gas corrosion resistance, if it is to meet utility boiler designers' requirements for 200,000 hours life at 700°C to 800°C. High temperature fireside corrosion in conventional boilers is caused by flue gas oxidation and liquid-phase coal ash corrosion. Coal ash corrosion is generally accepted as due to the presence of liquid sulfates on the surface of the tubing beneath an overlying ash deposit. The rate of this corrosion is alloy dependent and is also a function of the amount of sodium and potassium sulfate present in the coal ash and the level of SO_2 present in the flue gas. It is generally accepted that the severity of corrosion is at a maximum at approximately 700°C. Volatilization of the liquid sulfates above 700°C is responsible for reduction of the rate of corrosion at somewhat higher temperatures. To qualify the alloys for this service, screening tests were conducted at 700°C in a flue atmosphere of 15% CO_2-4%O_2-1.0%SO_2-Bal.N_2 flowing at the rate of 250 cubic centimeters per minute. The specimens were coated with a synthetic coal ash comprising 2.5% Na_2SO_4-2.5% K_2SO_4-31.67%Fe_2O_3-31.67%SiO_2-31.67%Al_2O_3. Pins were machined from vacuum melted, hot worked and annealed rod to approximately 9.5 mm diameter by 19.1 mm length. Each pin was given a 120 grit finish and subsequently coated using a water slurry of the coal ash. The weight of the coal ash coating was approximately 15 mg/cm². The screening was conducted for 1,000 hours after which the specimens were metallographically sectioned and the rate of metal loss and depth of attack by sulfidation determined. Specimens that exhibited a rate of metal loss or depth of sulfidation of less than 0.01 mm in 1,000 hours would have a corrosion loss of less than 2 mm in 200,000 hours. Table 3 presents these results for the compositions of Tables 2A and B.

TABLE 3

ALLOY	EQUATION I	CORROSION RATE (mm)	ALLOY	EQUATION I	CORROSION RATE (mm)
1	25.2	0.0026	A	21.6	0.0150
2	25.3	0.0010	B	22.3	0.0540
3	24.7	0.0100	C	18.1	0.0710
4	24.5	0.0050	D	30.7	0.0010
5	24.6	0.0100	E	17.2	0.0480
6	25.7	0.0076	G	30.0	0.0005
7	25.7	0.0064	H	22.8	0.0160
			I	48.2	0.0007
			J	22.1	0.0070
			K	32.2	0.0010
			L	24.1	0.0100
			M	29.0	0.0010

The alloys of Table 3 must pass the corrosion screen test of 0.01 mm or less corrosion

5   rate after 1,000 hours in coal ash/flue gas at 700°C for consideration of mechanical properties.

SCREENING FOR 100,000 HOURS/750°C/100 MPa STRESS RUPTURE LIFE

The utility boiler designers have determined that an alloy tubing capable of achieving

10   100,000 hours at 750°C and 100 MPa would meet their design criteria. This strength level
exceeds that of convention boiler tube alloys and virtually all iron- and nickel-base solid solution
alloys. The gamma prime strengthened alloys have the potential to achieve these strength levels.
However, these alloys typically possess low chromium levels which limit their corrosion
resistance and, for the most part, are too hard and low in ductility even in the annealed condition

15   to be fabricable into tubing. Alloy A in Table II is a gamma prime strengthened alloy that is
capable of both being fabricated into tubing and achieving the strength target. Referring to
Figure 1, alloys with data points plotted to the right of alloy A would also meet the strength
requirements, while those to the left of alloy A would fail prematurely at a temperature of 750°C
with a stress of 100 MPa for 100,000 hours. Table 4A contains the compositional limit value

20   and the pass/fail analysis of the alloys within the patent application and Table 4B presents the
results for the alloys of Table 2B. Note that alloys D, G, I, K, L and M which passed the
corrosion screen fail to meet the strength target; alloys A,B and C failed the corrosion screen but
passed the strength screen; while alloys E, F, H and J failed both screening tests.

TABLE 4A

ALLOY	EQUATION 3	STATUS
1	2.32	PASS
2	2.61	PASS
3	2.09	PASS
4	2.14	PASS
5	2.27	PASS
6	2.50	PASS
7	2.31	PASS

TABLE 4B

ALLOY	EQUATION 3	STATUS
A	2.69	PASS
B	2.90	PASS
C	5.37	PASS
D	2.85	FAIL
E	2.36	FAIL
F	2.45	FAIL
G	0.00	FAIL
H	1.38	FAIL
I	0.09	FAIL
J	0.20	FAIL
K	3.75	FAIL
L	1.63	FAIL
M	1.07	FAIL

Analysis of the microstructural phases using ThermoCalc (an analytical program that predicts phase stability based on thermodynamic data) lead to the conclusion that a minimum volume percent of 12% gamma prime was required to achieve the target stress rupture life of 750°C/100 MPa/100,000 hours and that the alloys containing approximately 18 to 20% gamma prime were too hard or too low in ductility to fabricate into tubing. Referring to Figure 2, the preferred compositional range when the titanium to niobium ratio is 1.5 to 0.7 is shown as a cross-hatched area. Four heats from Table 2 that are defined by this ratio are plotted by heat number.

SCREENING FOR MECHANICAL PROPERTY STABILITY

Mechanical property stability over the desired service life of 200,000 hours at the projected operating temperatures necessitates phase stability of the tubing alloy in order to withstand the stresses of periodic shutdowns, temperature fluctuations and hot spot variations over short lengths. High temperature alloys are especially susceptible to precipitation of carbide and embrittling phases in the temperature range of 700°C to 800°C. Limiting carbon to 0.3% and preferably less than 0.15% is essential if $Cr_{23}C_6$ and $(Cr,Mo)_6C$ type carbides are to be held

to acceptable levels. This is especially true because of the 22.6-30% chromium content required for corrosion resistance. Limiting the maximum content of chromium, molybdenum, aluminum, titanium and niobium elemental levels is required if embrittling phases such as sigma, mu and alpha-chromium are to be avoided. Table 5A presents the compositional limits as defined by

5   Equation (2) for the upper limits desired for maximum embrittling phase content for the alloys of this invention and Table 5B for the alloys outside the limits of this patent application for general information.

TABLE 5A

ALLOY	EQUATION 2
1	29.9
2	30.0
3	27.2
4	27.0
5	27.1
6	28.2
7	28.2

10

TABLE 5B

ALLOY	EQUATION 2
A	26.2
B	22.3
C	22.1
D	30.7
E	17.2
F	23.7
G	30.0
H	30.0
I	48.2
J	24.5
K	32.4
L	26.5
M	32.3

15   Of paramount importance is that the alloys be as free of sigma, mu and alpha-chromium as possible. That the alloys of this patent application were developed with this limitation in mind as shown in Figure 3. That the aluminum content is important is made clear by the role that aluminum plays in forming gamma prime (Ni_3Al), which diminishes the contribution that nickel exhibits in stabilizing the matrix against chromium-based phases. Selected alloys of Table 2A

20   are shown by their number designation as a function of their chromium and molybdenum contents. Surprisingly, this boundary is important with respect to molybdenum content in that alloys with excessive molybdenum (beyond its solubility limit) exhibited markedly decreased resistance to coal ash/flue gas corrosion. Figure 3 predicts alloys 3 and 4 would be completely

free of sigma phase while small amounts of sigma phase would potentially form in the remainder of the alloys of Table 2A. Table 6 contrasts the corrosion rate of the qualifying alloys, as a function of the molybdenum content, at a near constant chromium level of 23 weight percent.

TABLE 6

ALLOY	MOLYBDENUM CONTENT (%)	CHROMIUM CONTENT (%)	CORROSION RATE PER 1,000 H (MM)
3	3.09	22.93	0.0050
2	5.89	23.09	0.0026
H	9.00	22.00	0.0160

This nickel-base alloy range is useful for multiple-high-temperature-high-stress applications, such as, gas turbine engines and high-temperature boilers. The fabricability allows manufactures to form this alloy range into plate, sheet, strip or tubing with conventional fabricating equipment. The alloy range possesses the strength, corrosion resistance, thermal stability and manufacturability for advanced ultra-supercritical boiler tubing. Tubing of this alloy range having greater than 4 mm wall thickness with a 36 mm O.D. has the unique ability to exceed a stress rupture life of 100,000 hours at 100 MPa at 750°C and coal ash/flue gas corrosion of less than 2 mm for 200,000 hours at 700°C.

In accordance with the provisions of the statute, this specification illustrates and describes specific embodiments of the invention. Those skilled in the art will understand that the claims cover changes in the form of the invention and that certain features of the invention may operate advantageously without a corresponding use of the other features.

We claim:

1.   A nickel-base alloy consisting essentially of, by weight percent, about 10 to 24 cobalt, about 22.6 to 30 chromium, about 2.4 to 6 molybdenum, about 0 to 9 iron, about 0.2 to 3.2 aluminum, about 0.2 to 2.8 titanium, about 0.1 to 2.5 niobium, about 0 to 2 manganese,

5   about 0 to 1 silicon, about 0.01 to 0.3 zirconium, about 0.001 to 0.01 boron, about 0.005 to 0.3 carbon, about 0 to 4 tungsten, about 0 to 1 tantalum and balance nickel and incidental impurities, the alloy being further characterized by satisfying:

1) $\%Cr + 0.6 \times \%Ti + 0.5\% \times \%Al + 0.3 \times \%Nb \geq 24\%$;

10

2) $\%(Cr + 0.8\% \times Mo) + 0.6 \times \%Ti + 0.5 \times \%Al + 0.3 \times \%Nb \leq 37.5\%$;

3) $\%Al + 0.56 \times \%Ti + 0.29x \%Nb \geq 1.7\%$; and

15

4) $\%Al + 0.56 \times \%Ti + 0.29 \times \%Nb \leq 3.4\%$.

2.   The alloy of claim 1 containing about 12 to 23 cobalt, about 22.8 to 29 chromium, about 2.4 to 5 molybdenum, about 0.3 to 2.5 aluminum, about 0.3 to 2.7 titanium

20   and about 0.2 to 2.2 niobium.

3.   The alloy of claim 1 containing about 0.1 to 4 iron, about 0.1 to 1 manganese, about 0.1 to 0.8 silicon, about 0.01 to 0.2 zirconium, about 0.002 to 0.009 boron, about 0.01 to 0.2 carbon, about 0 to 2 tungsten and about 0 to 0.8 tantalum.

25

4.   The alloy of claim 1 containing about 12 to 20 volume percent gamma prime phase.

5.   The alloy of claim 1 characterized by a 100,000 hour 100 MPa stress rupture life

30   at a temperature of 750°C.

6.   A nickel-base alloy consisting essentially of, by weight percent, about 12 to 23 cobalt, about 22.8 to 29 chromium, about 2.4 to 5 molybdenum, about 0.1 to 4 iron, about 0.3 to 2.5 aluminum, about 0.3 to 2.7 titanium, about 0.2 to 2.2 niobium, about 0.1 to 1 manganese,

35   about 0.1 to 0.8 silicon, about 0.01 to 0.2 zirconium, about 0.002 to 0.009 boron, about 0.01 to

0.2 carbon, about 0 to 2 tungsten, about 0 to 0.8 tantalum and balance nickel and incidental impurities, the alloy being further characterized by satisfying:

1) $\%Cr + 0.6 \times \%Ti + 0.5\% \times \%Al + 0.3 \times \%Nb \geq 24\%$;

2) $\%(Cr + 0.8\% \times Mo) + 0.6 \times \%Ti + 0.5 \times \%Al + 0.3 \times \%Nb \leq 37.5\%$;

3) $\%Al + 0.56 \times \%Ti + 0.29x \%Nb \geq 1.7\%$; and

4) $\%Al + 0.56 \times \%Ti + 0.29 \times \%Nb \leq 3.4\%$.

7.   The alloy of claim 6 containing about 15 to 22 cobalt, about 23 to 28 chromium, about 2.5 to 3.5 molybdenum, about 0.5 to 2 aluminum, about 0.4 to 2.6 titanium and about 0.8 to 2.1 niobium.

8.   The alloy of claim 6 containing about 0.3 to 2 iron, about 0.1 to 0.5 manganese, about 0.2 to 0.7 silicon, about 0.03 to 0.15 zirconium, about 0.003 to 0.008 boron, about 0.02 to 0.15 carbon, about 0 to 1 tungsten and about 0 to 0.7 tantalum.

9.   The alloy of claim 6 containing about 12 to 20 volume percent gamma prime phase.

10.   The alloy of claim 6 characterized by a 100,000 hour 100 MPa stress rupture life at a temperature of 750°C.

11.   A nickel-base alloy consisting essentially of, by weight percent, about 15 to 22 cobalt, about 23 to 28 chromium, about 2.5 to 3.5 molybdenum, about 0.3 to 2 iron, about 0.5 to 2 aluminum, about 0.4 to 2.6 titanium, about 0.8 to 2.1 niobium, about 0.1 to 0.5 manganese, about 0.2 to 0.7 silicon, about 0.03 to 0.15 zirconium, about 0.003 to 0.008 boron, about 0.02 to 0.15 carbon, about 0 to 1 tungsten, about 0 to 0.7 tantalum and balance nickel and incidental impurities, the alloy being further characterized by satisfying:

1) $\%Cr + 0.6 \times \%Ti + 0.5\% \times \%Al + 0.3 \times \%Nb \geq 24\%$;

2) $\%(Cr + 0.8\% \times Mo) + 0.6 \times \%Ti + 0.5 \times \%Al + 0.3 \times \%Nb \leq 37.5\%$;

3)   %Al + 0.56 x %Ti + 0.29x %Nb ≥ 1.7%; and

4)   %Al + 0.56 x %Ti + 0.29 x %Nb ≤ 3.4%.

12.   The alloy of claim 11 containing about 18 to 21 cobalt, about 23.5 to 25 chromium, about 2.8 to 3.2 molybdenum, about 0.8 to 1.8 aluminum, about 0.5 to 2.5 titanium and about 1.2 to 2 niobium.

13.   The alloy of claim 12 containing about 0.5 to 1 iron, about 0.2 to 0.4 manganese, about 0.3 to 0.6 silicon, about 0.05 to 0.1 zirconium, about 0.004 to 0.007 boron, about 0.03 to 0.12 carbon, about 0 to 0.8 tungsten and about 0 to 0.5 tantalum.

14.   The alloy of claim 11 containing about 12 to 20 volume percent gamma prime phase.

15.   The alloy of claim 11 characterized by a 100,000 hour 100 MPa stress rupture life at a temperature of 750°C.

16.   The alloy of claim 11 wherein results of: equation 1) is at least 25%; equation 2) is less than or equal to 35.0%; equation 3) is at least 2.0%; and equation 4) is less than or equal to 3.0%.
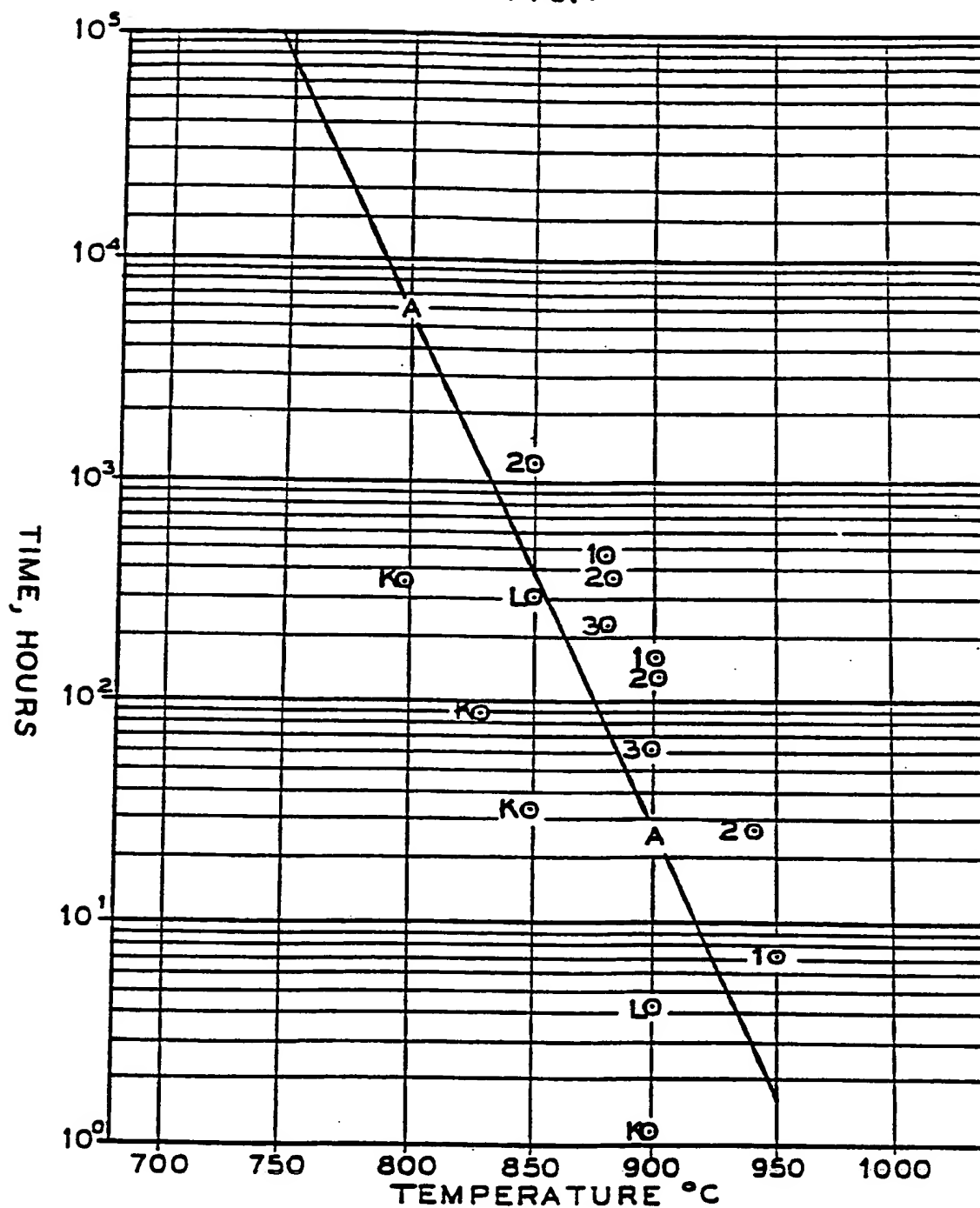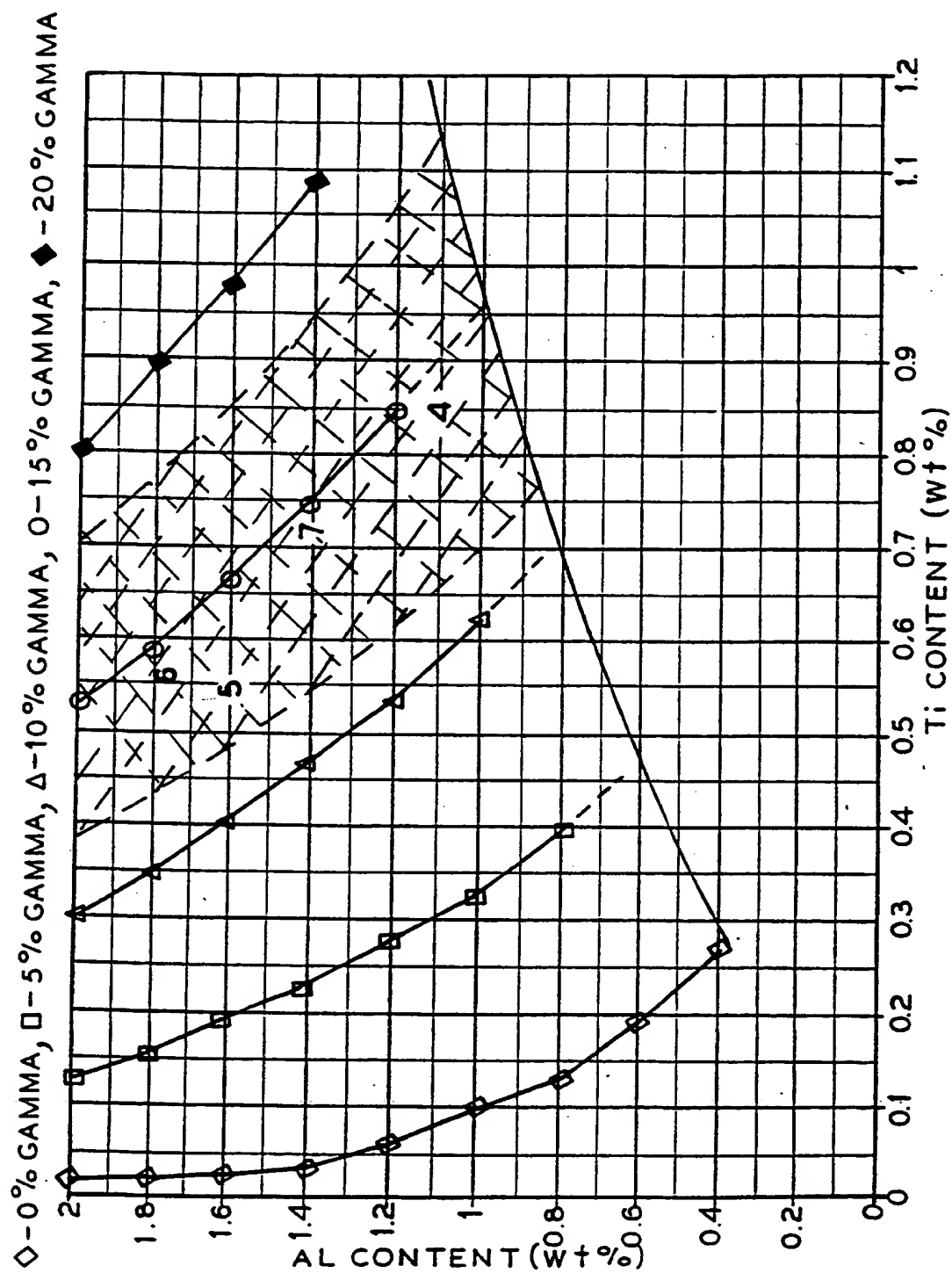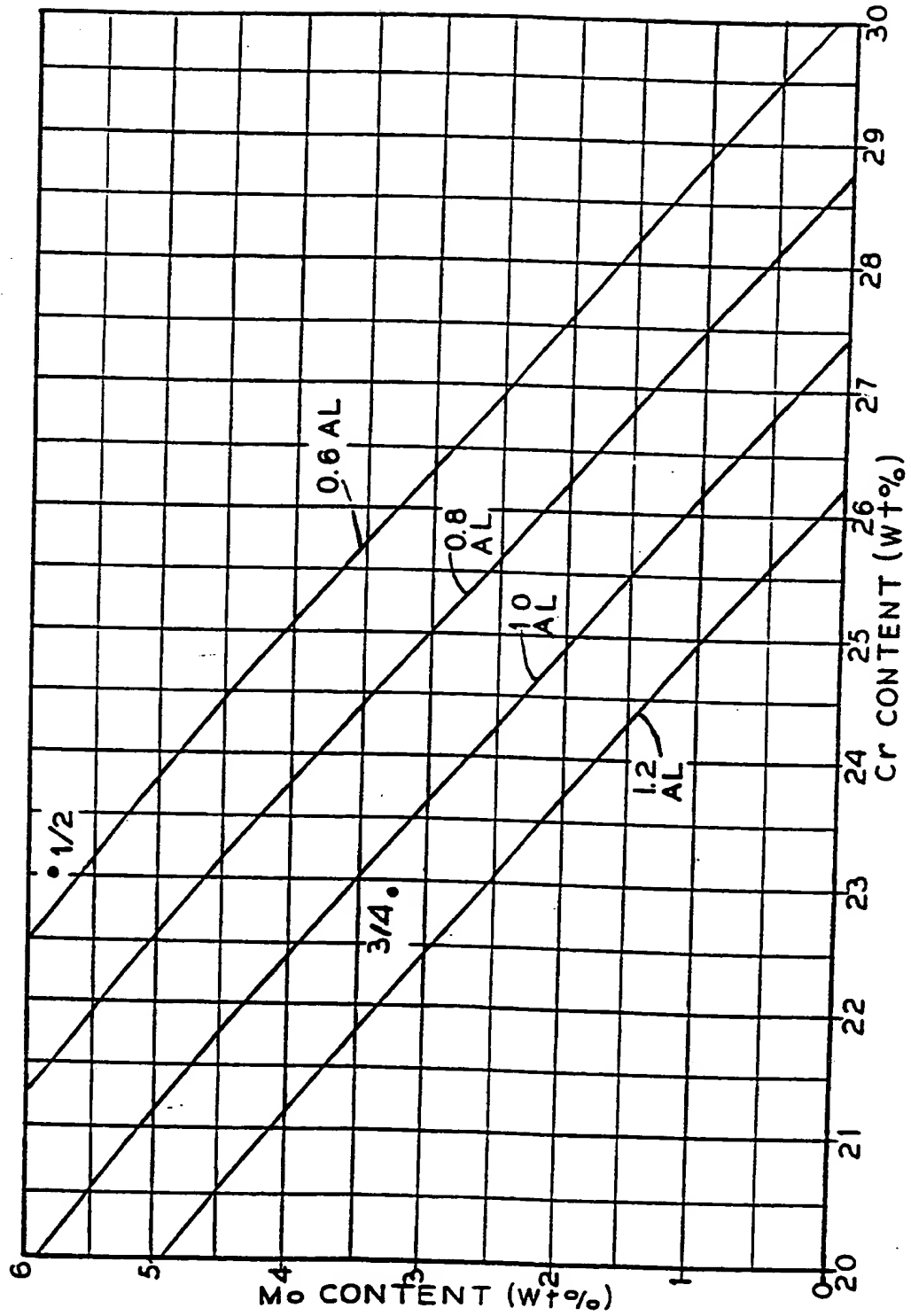
FIG.1

FIG.2

FIG.3

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US99/13059

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C22C 19/05
US CL :420/448

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 420/448, 588; 148/410, 428, 442

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

Please See Extra Sheet.

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 3,723,107 A (RICHARDS et al) 27 March 1973, col. 2, lines 28-60; col. 3-4, Table I; claim 1.	1-11 and 13-16
Y	US 4,039,330 A (SHAW et al.) 2 August 1977, col. 1, lines 22-26, 45-59, and 66; col. 2, lines 52-53; col. 4, lines 54-62; claim 1.	1-12 and 14-16

☐ Further documents are listed in the continuation of Box C.   ☐ See patent family annex.

*	Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance		
"E"	earlier document published on or after the international filing date	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O"	document referring to an oral disclosure, use, exhibition or other means		
"P"	document published prior to the international filing date but later than the priority date claimed	"&"	document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
12 NOVEMBER 1999	02 DEC 1999

Name and mailing address of the ISA/US	Authorized officer
Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231	ANDREW L OLTMANS
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B. FIELDS SEARCHED ·
Electronic data bases consulted (Name of data base and where practicable terms used):

WEST
search terms: nickel based, nickel-based, superalloy, super alloy, nickalloy, co or cobalt, cr or chromium, mo or molybdenum, al or aluminum, ti or titanium, nb or niobium or cb or columbium, zr or zirconium, carbon, boron, gamma prime phase, stress rupture life, fe or iron, si or silicon or tungsten or wolfram or ta or tantalum

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